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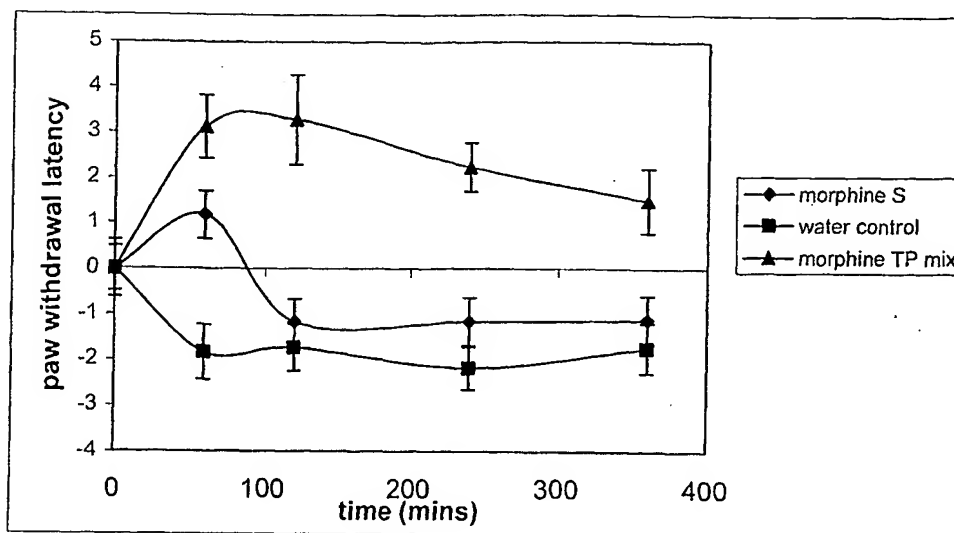
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(54) Title: CARRIER FOR ENTERAL ADMINISTRATION



(57) Abstract: There is provided a carrier for use in enteral administration of biologically active compounds, said carrier comprising an effective amount of one or more phosphate derivatives of one or more electron transfer agents.

## CARRIER FOR ENTERAL ADMINISTRATION

### Field of the Invention

- 5 This invention relates to a carrier for use in the enteral administration of biologically active compounds.

### Background of the Invention

- In this specification, where a document, act or item of knowledge is referred to or discussed, this reference or discussion is not to be taken as an admission that the document, act or item of
- 10 knowledge was at the priority date: part of common general knowledge; or known to be relevant to an attempt to solve any problem with which this specification is concerned.

- The major objective in drug delivery is to obtain an appropriate biological effect at a desired site of action. The choice of formulation can be critical to the efficacy of a drug since the bioactivity of a drug will be sub-optimal if it does not possess the correct physiochemical
- 15 properties to allow release from the formulation at the target site of action.

- Drugs are presented in different forms which are usually classified as enteral or parenteral. Enteral refers to forms of administration which involve entering the body through the gastrointestinal (*GI*) tract. Parenteral refers to all forms of administration which do not involve the GI tract. This classification highlights the fact that the GI tract introduces specific
- 20 pharmacokinetic issues which do not arise with parenteral routes, in particular, first pass metabolism through the liver. Generally the form of administration differs markedly as illustrated in the following table.

Enteral Administration

Pills, tablets and capsules

Enteric-coated preparations

Sustained-release preparations

Oral liquid preparations

Buccal / Sublingual administration

Rectal administration

Parenteral Administration

Intranasal administration

Subcutaneous administration

Intramuscular injections

Intravenous injections

Topical preparations

Transdermal administration

Enteral delivery involves administering the drug via the GI tract where the drug is absorbed and distributed via the bloodstream to the target site of action. Drugs delivered orally are  
5 absorbed through the intestine, drugs delivered buccally are absorbed through the mouth and drugs delivered rectally are absorbed through the rectum.

Oral delivery is dependent on the normal digestive processes of the GI tract. Once the oral drug formulation has been swallowed it travels through the esophagus to the stomach. The stomach has three tasks to do: (1) storage which requires the upper part of the stomach to relax  
10 and accept large volumes of swallowed material; (2) combine the swallowed material with digestive juice which requires the lower part of the stomach to mix these materials by its muscle action; and (3) empty its contents slowly into the small intestine. Several factors affect emptying of the stomach, including the nature of the food (mainly its fat and protein content) and the degree of muscle action of the emptying stomach and the small intestine. As the food  
15 is digested in the small intestine and dissolved into the juices from the pancreas, liver, and intestine, the contents of the intestine are mixed and pushed forward to allow further digestion. Finally, all of the digested nutrients are absorbed through the intestinal walls. The undigested materials are propelled into the colon and expelled. The normal digestive processes which control the rate of drug delivery to / removal from the target site are (1) when the stomach

empties the drug into the small intestine and (2) the time spent in the intestinal tract before absorption. Unfortunately, these processes cannot be controlled. However, by instructing a patient to take a drug before, after or with food it is possible to optimise the effect of these processes.

5 The chemical environment of the GI tract is also important to oral drug delivery. The drug must be in a form which is stable at the different pH of the various parts of the GI tract. If the drug forms a non-absorbable complex or is degraded chemically or enzymatically then this will decrease absorption. The drug must also be in solution in the GI fluids to be absorbed.

Sedimentation of the drug involves the drug forming solid particles and thus leaving the  
10 solution. Adsorption onto luminal solid particles involves solids adsorbing the drug, that is removing the drug from solution. Both sedimentation and adsorption decrease absorption of the drug. In many cases, degradation and complexation can be circumvented, or at least minimized, by chemical or formulation approaches so that they do not present a limitation to drug uptake.

15 Further, if a drug is absorbed through the intestinal or stomach wall, it then must pass through the liver. The liver is designed to eliminate foreign compounds from the body. As a result, a significant proportion of the drug (for example, 40-50%) may be metabolised and excreted before it reaches the bloodstream. It is possible to reduce the effect of the liver on enteral administration by having the drug absorbed through the lining of the mouth (buccal /  
20 sublingual) or the lining of the rectum (suppositories), however these routes are not always appropriate.

The role of the drug formulation in the delivery of drug to the target site of action should not be ignored. With any drug it is possible to alter its bioavailability considerably by formulation modification. Since a drug must be in solution to be absorbed from the GI tract, you may  
25 expect the bioavailability of a drug to decrease in the order: solution > suspension > capsule > tablet > coated tablet. This order may not always be followed but it is a useful guide.

Attempts to improve the bioavailability of enterally administered biologically active compounds involve either the formation of prodrugs, for example morphine sulphate or the use of excipients which improve absorption. There is still a need for enteral formulations which further improve the bioavailability of biologically active compounds.

## 5    **Summary of the Invention**

It has been found that a carrier composition comprising phosphates of electron transfer agents increases the efficacy of biologically active compounds administered enterally.

According to a first aspect of the invention, there is provided a carrier for use in enteral administration of biologically active compounds, said carrier comprising an effective amount  
10    of one or more phosphate derivatives of one or more electron transfer agents.

According to a second aspect of the invention, there is provided a method for improving the efficacy and transport of enterally administered biologically active compounds, said method comprising the step of combining the biologically active compound with a carrier comprising an effective amount of one or more phosphate derivatives of one or more electron transfer  
15    agents.

Preferably, the phosphate derivative of an electron transfer agent is selected from the group consisting of phosphate derivatives of tocopherol, phosphate derivatives of tocotrienol and mixtures thereof.

The present invention also provides for the use of an effective amount of one or more  
20    phosphate derivatives of one or more electron transfer agents, such as phosphate derivatives of tocopherol or tocotrienol, together with other excipients, in the manufacture of a carrier for use in the enteral administration of biologically active compounds.

The present invention also provides a pharmaceutical composition for enteral administration comprising one or more biologically active compounds and a carrier comprising an effective

amount of one or more phosphate derivatives of one or more electron transfer agents, such as phosphate derivatives of tocopherol or tocotrienol.

According to a third aspect of the invention, there is provided a method for improving the efficacy and transport of enterally administered biologically active compounds, said method comprising the step of combining the biologically active compound with a carrier comprising an effective amount of one or more complexes of phosphate derivatives of an electron transfer agent.

According to a fourth aspect of the invention, there is provided a carrier for use in enteral administration of biologically active compounds, said carrier comprising an effective amount of one or more complexes of phosphate derivatives of an electron transfer agent.

In one preferred form of the invention, the carrier and biologically active compound are in a form which is protected by an enteric coating. The enteric coating must be insoluble in the stomach (low pH) and survive the enzymes in the saliva, but degrade in the absorption site which is just after the stomach at a pH greater than 6. Typically, the coating is a water soluble polymer such as cellulose ether, polyvinylpyrrolidone or polyethylene glycol. The advantage of the enteric coating is that it increases the likelihood that the carrier and the biologically active compound will be in proximity to each other at the absorption site and thus maximises the effects of the carrier upon the efficacy and transport of the biologically active compound. For example, a tablet or capsule may have an enteric coating or a functional food may contain microencapsulated particles which have an enteric coating.

The term "carrier" is used in this specification to include all forms of enteral administration. It includes but is not limited to pills, tablets, capsules, liquid formulations, functional foods, dietary supplements, lozenges, suppositories.

The term "effective amount" is used herein to refer to an amount of the one or more phosphate derivatives of an electron transfer agent that assists the biologically active compound to be

absorbed in an amount that is measurably effective in the reduction of one or more symptoms presented by a patient. The effective amount may range up to 99.99% w/w of the total weight of the carrier. A person skilled in the art will understand that the actual amount will vary depending upon the biologically active compound. The effective amount will be sufficient to

5 deliver an amount of biologically active compound within the therapeutic range of that biologically active compound. The effective amount used will also depend on whether the phosphate derivative of an electron transfer agent is being used to assist with formulation properties, for example, solubilisation or surface activity. Where the phosphate derivative of an electron transfer agent is acting as a solubiliser, the effective amount will depend on the

10 concentration of the drug in the formulation and may range from 40% to 90% w/w, preferably 45 to 75% w/w, more preferably 50 to 60% w/w. Where the phosphate derivative of an electron transfer agent is not required for solubilisation properties, the effective amount may be in the range of 0.01 to 20% w/w, preferably 1 to 15% w/w and more preferably 5 to 10% w/w. Preferably (when solubilisation properties are not required), the effective amount of the one or

15 more phosphate derivatives of an electron transfer agent is in the range of from 0.1 to 10 % w/w of the total weight of the carrier. More preferably, in the range of 5 to 10% and most preferably 7.5% w/w.

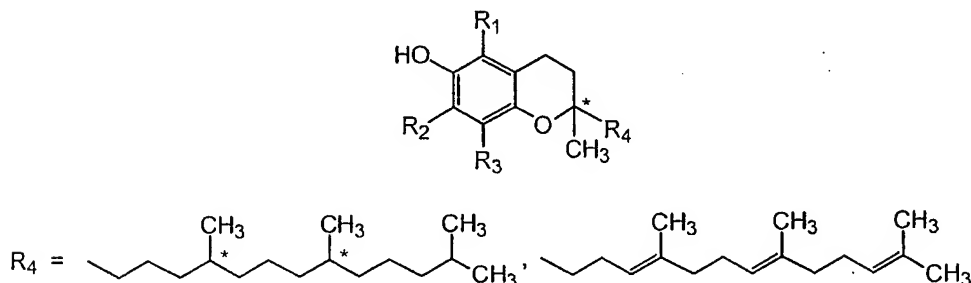
The term "electron transfer agents" is used herein to refer to the class of chemicals which may be phosphorylated and which (in the non-phosphorylated form) can accept an electron to

20 generate a relatively stable molecular radical or accept two electrons to allow the compound to participate in a reversible redox system. Examples of classes of electron transfer agent compounds that may be phosphorylated include hydroxy chromans including alpha, beta, gamma and delta tocopherols in enantiomeric and racemic forms; quinols being the reduced forms of electron transfer agent K1 and ubiquinone; hydroxy carotenoids including retinol; calciferol

25 and ascorbic acid. Preferably, the electron transfer agent is selected from the group consisting of tocopherol and other tocopherols, retinol, electron transfer agent K1 and mixtures thereof.

More preferably, the electron transfer agent is selected from the group consisting of the tocopherols and mixtures thereof. The tocopherols include all isomers of derivatives of 6-hydroxy 2-methyl chroman (see structure below) where  $R_1$ ,  $R_2$  and  $R_3$  may be hydrogen or methyl groups, that is, the  $\alpha$ -5:7:8 tri-methyl;  $\beta$ -5:8 di-methyl;  $\gamma$ -7:8 di-methyl; and  $\delta$  8 methyl derivatives. In the

5 tocopherols,  $R_4$  is substituted by 4:8:12 tri-methyl tridecane and the 2, 4, and 8 positions (see \*) may be stereoisomers with R or S activity or racemic. In the tocotrienols,  $R_4$  is substituted by 4:8:12 tri-methyl trideca-3:7:11 triene and the 2 position may be stereoactive as R or S stereoisomers or racemic. Most preferably, the electron transfer agent is  $\alpha$ -tocopherol or tocotrienol.



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The term "phosphate derivatives" is used herein to refer to the acid forms of phosphorylated electron transfer agents, salts of the phosphates including metal salts such as sodium, magnesium, potassium and calcium and any other derivative where the phosphate proton is replaced by other substituents such as ethyl or methyl groups or phosphatidyl groups. The

15 term includes mixtures of phosphate derivatives, especially those which result from phosphorylation reactions, as well as each of the phosphate derivatives alone. For example, the term includes a mixture of mono-tocopheryl phosphate (TP) and di-tocopheryl phosphate (T2P) as well as each of TP and T2P alone. Suitable mixtures are described in international patent application no PCT/AU01/01475.



Preferably, the one or more phosphate derivatives of one or more electron transfer agents is selected from the group consisting of mono-tocopheryl phosphate, di-tocopheryl phosphate, mono-tocotrienyl phosphate, di-tocotrienyl phosphate and mixtures thereof. Most preferably, the one or more phosphate derivatives of one or more electron transfer agents is a mixture of one or more of mono-tocopheryl phosphate, di-tocopheryl phosphate, mono-tocotrienyl phosphate and di-tocotrienyl phosphate.

In some situations, it may be necessary to use a phosphate derivative such as a phosphatide where additional properties such as increased water solubility are preferred. Phosphatidyl derivatives are amino alkyl derivatives of organic phosphates. These derivatives may be prepared from amines having a structure of  $R_1R_2N(CH_2)_nOH$  wherein  $n$  is an integer between 1 and 6 and  $R_1$  and  $R_2$  may be either H or short alkyl chains with 3 or less carbons.  $R_1$  and  $R_2$  may be the same or different. The phosphatidyl derivatives are prepared by displacing the hydroxyl proton of the electron transfer agent with a phosphate entity that is then reacted with an amine, such as ethanolamine or N,N' dimethylethanolamine, to generate the phosphatidyl derivative of the electron transfer agent. One method of preparation of the phosphatidyl derivatives uses a basic solvent such as pyridine or triethylamine with phosphorous oxychloride to prepare the intermediate which is then reacted with the hydroxy group of the amine to produce the corresponding phosphatidyl derivative, such as P cholyl P tocopheryl dihydrogen phosphate.

In some situations, complexes of phosphate derivatives of the electron transfer agents may also be utilized where additional properties such as improved stability or deliverability may be useful. The term "complexes of phosphate derivatives" refers to the reaction product of one or more phosphate derivatives of electron transfer agents with one or more complexing agents selected from the group consisting of amphoteric surfactants, cationic surfactants, amino acids having nitrogen functional groups and proteins rich in these amino acids as disclosed in international patent application no PCT/AU01/01476, incorporated herein by reference.

The preferred complexing agents are selected from the group consisting of arginine, lysine and tertiary substituted amines, such as those according to the following formula:



wherein  $\text{R}^1$  is chosen from the group comprising straight or branched chain mixed alkyl radicals from C6 to C22 and carbonyl derivatives thereof;

$\text{R}^2$  and  $\text{R}^3$  are chosen independently from the group comprising H,  $\text{CH}_2\text{COOX}$ ,  $\text{CH}_2\text{CHOHCH}_2\text{SO}_3\text{X}$ ,  $\text{CH}_2\text{CHOHCH}_2\text{OPO}_3\text{X}$ ,  $\text{CH}_2\text{CH}_2\text{COOX}$ ,  $\text{CH}_2\text{COOX}$ ,  $\text{CH}_2\text{CH}_2\text{CHOHCH}_2\text{SO}_3\text{X}$  or  $\text{CH}_2\text{CH}_2\text{CHOHCH}_2\text{OPO}_3\text{X}$  and X is H, Na, K or alkanolamine provided  $\text{R}^2$  and  $\text{R}^3$  are not both H; and

wherein when  $\text{R}^1$  is RCO then  $\text{R}^2$  may be  $\text{CH}_3$  and  $\text{R}^3$  may be  $(\text{CH}_2\text{CH}_2)\text{N}(\text{C}_2\text{H}_4\text{OH})\text{H}_2\text{CHOPO}_3$  or  $\text{R}^2$  and  $\text{R}^3$  together may be  $\text{N}(\text{CH}_2)_2\text{N}(\text{C}_2\text{H}_4\text{OH})\text{CH}_2\text{COO}^-$ .

Preferred complexing agents include arginine, lysine or lauryliminodipropionic acid where complexation occurs between the alkaline nitrogen centre and the phosphoric acid ester to form a stable complex.

The term "biologically active compound" is used herein to refer to compounds having a biological effect in humans or animals for medical or veterinary application. Biologically active compounds include pharmaceuticals, nutritional supplements, drugs, vitamins, phytochemicals, cosmeceuticals, nutraceuticals, nutrients and other health supplements that are useful for the treatment of human beings or other animals for prophylaxis or therapy.

Examples of biologically active compounds include but are not limited to narcotic analgesics such as morphine and levorphanol, non narcotic analgesics such as codeine and acetaminophen, corticosteroids such as cortisone, anaesthetics such as propofol, antiemetics such as scopolamine, sympathomimetic drugs such as adrenaline and dopamine, antiepileptic drugs such as fosphenytoin, anti-inflammatory drugs such as ibuprofen, thyroid hormones and antithyroid drugs including thyroxine, phytochemicals including  $\alpha$ -bisabolol, eugenol, silybin,

soy isoflavones, iridoid glycosides including aucubin and catalpol, sesquiterpene lactones including pseudoguaianolide from *Arnica chamissonis*, terpenes including rosmarinic acid and rosmanol, phenolic glycosides including the salicylates salicin, saligenin and salicyclic acid, triterpenes taxasterol or  $\alpha$ -lactuceryl, and isolactuceryl, *p*-hydroxyphenylacetic acid derivative

5 taraxacoside, hydroquinone derivatives including arbutin, phenylalkanones including gingerols and shagaols, hypericin, statins, and acylphloroglucides including xanthohumol, lupulone, humulone and 2-methylbut-3-en-2-ol. Examples of nutrients and nutraceuticals include vitamins, important precursor molecules for generation of hormones, minerals, proteins, minerals, amino acids, plant extracts such as grape seed extract, ephedrine, DHEA,

10 isoflavones, phytosterols and similar bioagents. The biologically active compound may also be a peptide, polypeptide or protein. The biologically active compound can be in any suitable form including phosphate derivatives.

A person skilled in the art would know which other excipients could be included in the carrier. The choice of other excipients would depend on the characteristics of the biologically active

15 compound. Examples of other excipients include solvents, surfactants, emollients, preservatives and the like. The choice of other excipients will also depend on the form of administration used.

#### **Brief Description of the Drawings**

Figure 1: Effect of morphine sulphate 5 mg/kg, morphine with tocopheryl phosphate carrier

20 according to the invention 5 mg/kg and control on paw withdrawal latency in rats, tested over 3 hours (pooled data).

Figure 2: CoQ<sub>10</sub> and CoQ<sub>9</sub> standard curves

## Examples

The invention is further explained and illustrated by the following non-limiting examples.

### Example 1

In this experiment, the efficacy of a morphine composition according to the invention was compared with the efficacy of morphine sulphate, the currently used enteral formulation of morphine. The effect was measured by comparison of times taken for a rat to withdraw its paw in response to heat when medicated and unmedicated with morphine.

### Materials

Animals: Nine conscious Sprague-Dawley rats weighing between 350-450 grams each

10 Treatment groups:

1. Control: water,
2. morphine sulphate,
3. Morphine with TPm: morphine HCl (14%) in a carrier containing water (59%) and a tocopheryl phosphate mixture (27%) (*TPm*). The TPm contained mono-tocopheryl phosphate and di-tocopheryl phosphate.

15

Formulations 2 and 3 were diluted with water and the final morphine concentration was made up to 5 mg/ml. For example, 0.357 grams of formulation 3 was mixed with 0.643 grams of water to obtain a final morphine concentration of 5%. This liquid formulation was then delivered to the animals by oral gavage (tube into stomach).

20 Method

The experiment used nine rats that were divided into three groups. After the first treatment, the rats were rested and each group was given a different treatment. The process was repeated once more until each rat had been given each of the three treatments.

Water, morphine sulphate and morphine with TPm were given by oral gavage at a concentration of 5mg/kg of body weight. Analgesic testing was performed at 1, 2, 4 and 6

25

hours and at each time point withdrawal latency was measured three times on each rat (with at least five minutes rest if using the same paw).

A plantar analgesiometer designed for rapid and efficient screening of analgesia levels in small laboratory animals was used. The device applied a heat source ( $\sim 45^{\circ}\text{C}$  from an infrared light) to the animal's hindpaw and the time taken to withdraw the paw from the heat source was measured (paw withdrawal latency). The heat source (plate) provided a constant surface temperature. It had a built-in digital thermometer with an accuracy of  $0.1^{\circ}\text{C}$  and a timer with an accuracy of 0.1 second. The animal was placed on a hot plate, confined by a clear acrylic cage which surrounds the plate and paw withdrawal response was monitored. An increased time period before paw withdrawal response indicating analgesia. Each animal was tested 3 times at each time point. (ie a single rat had the heat applied to its back foot three times at each time point).

The results are illustrated in Figure 1. Both the morphine sulphate and morphine in the tocopheryl phosphate carrier caused an increase in latency indicating analgesia. The morphine in the tocopheryl phosphate carrier caused a greater latency which was maintained for a longer period of time than the morphine sulphate. That is, the morphine formulated in a carrier according to the invention provided a sustained analgesic effect for up to 2 hours following oral administration whereas the morphine sulphate only provided an analgesic effect for the first hour. The standard error bars on the graph points do not overlap except at the one hour time point where the aqueous morphine sulphate and the morphine TPM formulation were similarly active. For the later time points, the morphine TPM formulation gave sustained analgesia.

1 Statistical Analysis: Comparison between morphine sulphate and Morphine TPM formulations.

25 • at 60mins  $t=2.598$  ( $p<0.02$ )

- at 120mins  $t=4.815$  ( $p<0.0005$ )
- at 240mins  $t=4.351$  ( $p<0.001$ )
- at 360 mins  $t=3.094$  ( $p=0.005$ )

### Conclusion

- 5 The use of the TPm carrier provided a sustained analgesia over a longer period of time using the same amount of morphine as the morphine sulphate formulation. Whilst the results were not significant at the one hour time point, the TPm formulation was statistically significant at all later time points.

### **Example 2**

- 10 This example investigates the bioavailability in guinea pigs of CoQ10 administered in the following formulations:

- A. CoQsol
- B. CoQsol plus TPM in MCT
- C. MCT oil (control)

15 Materials and Methods

#### *Formulations*

Tocopheryl phosphate mixture (TPM) containing monotocopheryl phosphate (TP) and ditocopheryl phosphate (T2P) in a ratio of 2:1 w/w was prepared by Phosphagenics Ltd.

CoQsol was purchased from Doctor's Trust Vitamins, U.S.A

- 20 Medium chain triglyceride (MCT) was manufactured by Abitec Corp, U.S.A.

The formulations consisted of the following:

A. CoQsol: Each softgel capsule contains 60 mg of CoQ, with the oily contents of the pills measuring 0.44 ml in volume. Therefore, the concentration of CoQ is  $60 \text{ mg}/0.44 \text{ ml} = 136 \text{ mg CoQ/ml}$  of capsule contents. Each millilitre of the CoQsol formulation also contains 136 IU d-

$\alpha$ -tocopherol and 3705 IU vitamin A. Excipients are rice bran oil, gelatin, glycerin, water, beeswax, annato extract and titanium dioxide.

B. CoQsol+TPM: the formulation was prepared with the concentration of CoQ such that 30 mg/kg was administered in the same volume relative to body weight of the treatment group; i.e., in volumes of approximately 0.21 ml per kg b.wt. Each ml of formulation contained CoQ and TPM, each at 140 mg/ml, with MCT as the diluent.

C. MCT: (vehicle): the control group received MCT at 0.21 ml/kg b.wt.

### *Animals*

Adult female guinea pigs were purchased from Animal Services, Monash University and acclimatised to the Departmental Animal House for a minimum of 5 days before the treatments commenced. Animals were randomly assigned to treatment groups (n=10), tagged with unique identifying marks on their backs (clipped hair and colour codes), and housed as a group in an environmentally-enriched pen of approximately 1 x 4 m in size. The average body weight of the CoQsol-treated group was 0.795 kg at day 0. The average body weight of the CoQsol+TPM group was 0.746 kg at day 0. The average body weight of the control group was 0.796kg at day 0.

Food and water: Rabbit and guinea pig standard laboratory pellets (Barastoc, Australia). Water was provided freely

Route and method of dosing: animals were dosed by oral gavage using a plastic cannula attached to a delivery syringe in volumes of approximately 0.21 ml per kg body weight.

### *Methods*

Dose of CoQsol: 30 mg / kg body weight / day

Dose of TPM: 30 mg / kg body weight / day

Dosing regimen: once daily

Dosing period: 26 days

Body weight was measured weekly.

At the completion of the treatment period, the guinea pigs were killed by asphyxiation using CO<sub>2</sub> gas. The blood was removed by heart puncture into heparinised collection tubes, and centrifuged for separation of plasma and stored at -80°C until extraction of CoQ.

- 5 Extractions of CoQ and analyses by HPLC were performed essentially according to the method of Aberg *et al.*, (1992) "Distribution and redox state of ubiquinones in rat and human tissues" *Arch. Biochem. Biophys.* 295: 230-34.

Increased levels of CoQ10 and CoQ9 are indicative of increased bioavailability and uptake.

- Both CoQ9 and CoQ10 were measured because guinea pigs can synthesize both forms of CoQ  
10 (9 and 10). It is therefore important to assess the levels of both forms following administration since administration of CoQ10 can increase both levels in vivo.

#### Results:

CoQ<sub>10</sub> and CoQ<sub>9</sub> concentration in plasma

PLASMA	CoQ <sub>9</sub> Peak area	CoQ <sub>10</sub> Peak area	CoQ <sub>9</sub> Conc (ng/ml)	CoQ <sub>10</sub> Conc (ng/ml)	CoQ <sub>9</sub> Conc (ng/vial)	CoQ <sub>10</sub> Conc (ng/vial)
Control						
PlasmaC1	0.191146	3.89095	22	292	11	146
PlasmaC2	1.26553	2.62247	176	197	88	98
PlasmaC3	0.69327	1.15616	94	87	47	43
PlasmaC4	0.874554	2.29567	120	172	60	86
PlasmaC5	0.558185	2.45003	74	184	37	92
PlasmaC6	0.90064	5.16449	123	388	62	194
PlasmaC7	1.0674	3.34757	147	251	74	126
PlasmaC8	0.190272	3.20794	22	241	11	120
PlasmaC9		3.18797		239		120
PlasmaC10		1.14622		86		43
Mean					49	107



	CoQ <sub>9</sub> Peak area	CoQ <sub>10</sub> Peak area	CoQ <sub>9</sub> Conc (ng/ml)	CoQ <sub>10</sub> Conc (ng/ml)	CoQ <sub>9</sub> Conc (ng/vial)	CoQ <sub>10</sub> Conc (ng/vial)
<b>PLASMA</b>						
<b>CoQSol</b>						
PlasmaS1	0.902389	5.18353	124	389	62	195
PlasmaS2	0.19846	2.45755	23	184	12	92
PlasmaS3	0.32906	2.60398	42	195	21	98
PlasmaS4	0.45307	3.0014	59	225	30	113
PlasmaS5	0.53315	2.3862	71	179	35	90
PlasmaS6	0.93297	2.04795	128	154	64	77
PlasmaS7	0.54571	2.1518	73	161	36	81
PlasmaS8	1.41097	2.16559	196	162	98	81
PlasmaS9	0.690034	6.3554	93	477	47	239
PlasmaS10	0.28032	3.61253	35	271	17	136
Mean					42	120
<b>CoQsol+TPM</b>						
PlasmaT1	0.84019	2.75545	115	207	57	103
PlasmaT2		4.47825		336		168
PlasmaT3	0.17022	4.3493	19	327	10	163
PlasmaT4	0.503096	4.92475	67	370	33	185
PlasmaT5	0.372346	3.01239	48	226	24	113
PlasmaT6	0.267389	6.29373	33	473	16	236
PlasmaT7		5.53598		416		208
PlasmaT8	0.26282	5.51973	32	415	16	207
PlasmaT9	0.336923	4.1363	43	311	21	155
PlasmaT10	0.105107	3.49962	10	263	5	131
Mean					23	167

The above results do not include any statistical information because as is known in the area of micronutrient studies, there is always a significant inter-animal variation due to the fact that each animal has different needs for CoQ10 and it will only be absorbed when needed. The

large inter-animal variation leads to large standard deviation figures, which do not accurately reflect the significance of the results.

Figure 2 shows the standard curves obtained in the analysis of CoQ10 and CoQ9 in the plasma before running the samples on the HPLC (the closer  $r^2$  is to 1 the more accurate the results).

5    Discussion

The results in Figure 2 and the above tables show that the CoQ levels in plasma are higher in CoQsol+TPM treated animals than in the animals treated with CoQsol alone or the control.

This illustrates that the tocopheryl phosphate mixture increases the bioavailability of CoQ10.

10    The word 'comprising' and forms of the word 'comprising' as used in this description and in the claims does not limit the invention claimed to exclude any variants or additions.

Modifications and improvements to the invention will be readily apparent to those skilled in the art. Such modifications and improvements are intended to be within the scope of this invention.

**The claims defining the invention are as follows:**

- 1 A method for improving the efficacy and transport of enterally administered biologically active compounds, said method comprising the step of combining the biologically active compound with a carrier comprising an effective amount of one or more phosphate derivatives of one or more electron transfer agents.
- 2 The method according to claim 1 wherein the electron transfer agent is selected from the group consisting of hydroxy chromans including alpha, beta, gamma and delta tocopherols in enantiomeric and racemic forms; quinols being the reduced forms of electron transfer agent K1 and ubiquinol; hydroxy carotenoids including retinol; calciferol and ascorbic acid.
- 3 The method according to claim 2 wherein the electron transfer agent is selected from the group consisting of tocopherol and other tocopherols, retinol, electron transfer agent K1 and mixtures thereof.
- 4 The method according to claim 3 wherein the electron transfer agent is selected from the group consisting of tocopherols and mixtures thereof.
- 5 The method according to claim 4 wherein the phosphate derivative of an electron transfer agent is selected from the group consisting of phosphate derivatives of tocopherol, phosphate derivatives of tocotrienol and mixtures thereof.
- 6 The method according to claim 5 wherein the phosphate derivative of an electron transfer agent is selected from the group consisting of mono-tocopheryl phosphate, di-tocopheryl phosphate, mono-tocotrienyl phosphate, di-tocotrienyl phosphate and mixtures thereof.
- 7 The method according to claim 1 wherein the phosphate derivative is selected from the group consisting of phosphates, phosphatides, complexes of phosphates and mixtures thereof.

- 8 The method according to claim 1 further comprising the step of protecting the combined carrier and biologically active compound with an enteric coating.
- 9 The method according to claim 1 wherein the biologically active compound is morphine.
- 10 The method according to claim 1 wherein the biologically active compound is co-enzyme Q10.
- 11 A carrier for use in enteral administration of biologically active compounds, said carrier comprising an effective amount of one or more phosphate derivatives of one or more electron transfer agents.
- 12 Use of an effective amount of one or more phosphate derivatives of one or more electron transfer agents, together with other excipients, in the manufacture of a carrier for use in the enteral administration of biologically active compounds.
- 13 A pharmaceutical composition for enteral administration comprising one or more biologically active compounds and a carrier comprising an effective amount of one or more phosphate derivatives of one or more electron transfer agents.
- 14 A method for improving the efficacy and transport of enterally administered biologically active compounds, said method comprising the step of combining the biologically active compound with a carrier comprising an effective amount of one or more complexes of phosphate derivatives of an electron transfer agent.
- 15 A carrier for use in enteral administration of biologically active compounds, said carrier comprising an effective amount of one or more complexes of phosphate derivatives of an electron transfer agent.

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Figure 1

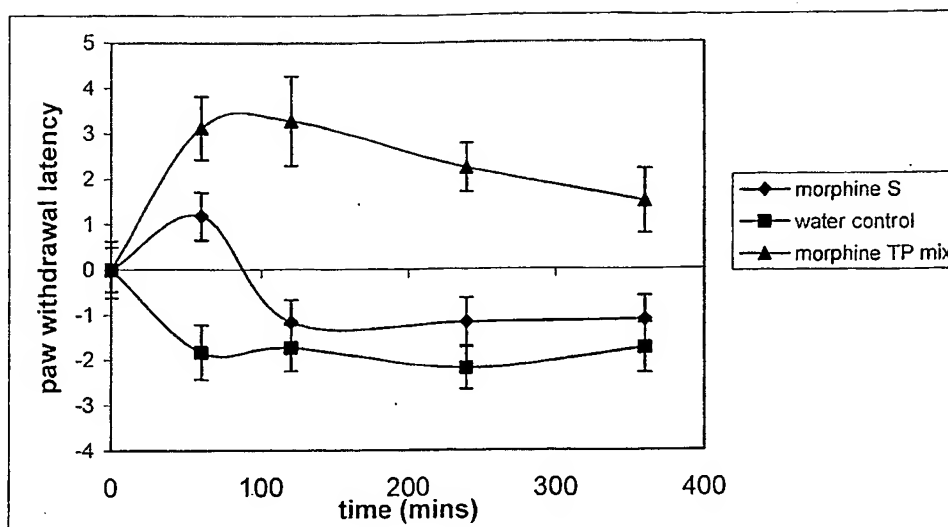
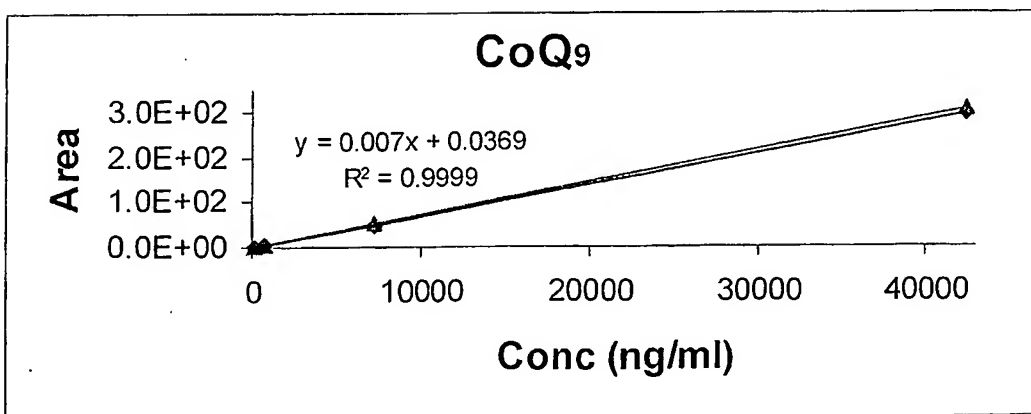
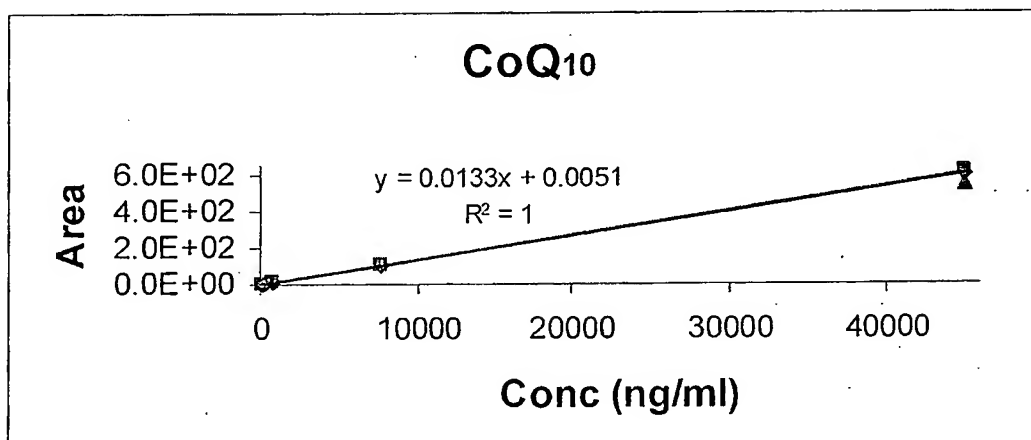


Figure 2



# INTERNATIONAL SEARCH REPORT

International application No.  
**PCT/AU2005/001159**

**A. CLASSIFICATION OF SUBJECT MATTER**

Int. Cl. <sup>7</sup>: A61K 47/22, 31/355, 31/375, 31/07, 31/59, A61P 29/00

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

DWPI: tocopherol, tocotrienol, tocol, chroman, quinol, ubiquinone, carotenoid, retinol, calciferol, ascorbic acid, phosphate, tocopherol phosphate, tocopheryl phosphate, and enteral.

CA: As above

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2002/040034 A1 (Tocovite Pty Ltd) 23 May 2002, See whole document.	1-15
X	WO 2002/040033 A1 (Tocovite Pty Ltd) 23 May 2002, See whole document.	1-15
X	WO 2003/011303 A1 (Vital Health Sciences Pty Ltd) 13 February 2003, See whole document.	11, 13, 15
X	WO 2004/014432 A1 (Vital Health Sciences Pty Ltd) 19 February 2004, See whole document.	11, 13, 15



Further documents are listed in the continuation of Box C



See patent family annex

* Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search  
**31 August 2005**

Date of mailing of the international search report  
**08 SEP 2005**

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# INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/AU2005/001159

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report			Patent Family Member		
WO 0240034	AU	14820/02	AU	14821/02	BR 0115952
	BR	0115953	CA	2426852	CA 2426885
	CN	1474697	CN	1474698	EP 1339412
	EP	1339413	MX	PA03003584	MX PA03003585
	US	2004052754	US	2004096493	US 2004097472
	WO	0240033			
WO 03011303	BR	0211673	CA	2453823	CN 1547475
	EP	1420797	MX	PA04000654	US 2004253318
WO 2004014432	AU	2003246461	BR	0313575	CA 2495280
	EP	1545621			
Due to data integration issues this family listing may not include 10 digit Australian applications filed since May 2001.					
END OF ANNEX					